-continued

Components	Composition Per Unit (mg)
Shell	
Iron oxide, red (C.I. No. 77491, EC No. E 172)	0.123
Iron oxide, yellow (C.I. No. 77492, EC No. E 172)	0.123
Iron oxide, black (C.I. No. 77499, EC No. E 172)	0.245
Titanium dioxide	1.540
Gelatin	74.969
Total Tablet Mass	342.00

The formulation is manufactured, e.g., as described in Formulation Example 4.

Formulation Example 6

Hard Gelatine Capsule

Components	Composition Per Unit (mg)
Valsartan [= active ingredient]	80.00
Sodium lauryl sulphate	0.60
Magnesium stearate	1.30
Povidone	12.50
Crospovidone	13.00
Microcystalline cellulose	21.10
Total Tablet Mass	130.00

Formulation Example 7

A hard gelatin capsule, comprising as active ingredient, e.g., (S)—N-(1-carboxy-2-methylprop-1-yl)-N-pentanoyl-N-[2'(1H-tetrazol-5-yl)biphenyl-4-yl-methyl]amine, can be formulated, e.g., as follows:

Components	Composition Per Unit (mg)
(1) Valsartan	80.00
(2) Microcystalline cellulose	110.0
(3) Polyvidone K30	45.2
(4) Sodium lauryl sulfate	1.2

-continued

Components	Composition Per Unit (mg)
(5) Crospovidone	26.0
(6) Magnesium stearate	2.6

Components (1) and (2) are granulated with a solution of components (3) and (4) in water. The components (5) and (6) are added to the dry granulate and the mixture is filled into size 1 hard gelatin capsules.

All publications and patents mentioned herein are incorporate by reference in their entirety as if set forth in full herein.

What is claimed is:

1. A method for the treatment of a condition or disease selected from the group consisting of hypertension and heart failure, comprising administering to a patient in need thereof a therapeutically effective amount of the combination of:

(i) the AT 1-antagonist valsartan or a pharmaceutically acceptable salt thereof;

(ii) the NEP inhibitor N-(3-carboxy-1-oxopropyl)-(4S)-(p-phenylphenylmethyl)-4-amino-2R-methylbutanoic acid ethyl ester or a pharmaceutically acceptable salt thereof, or (2R,4S)-5-biphenyl-4-yl-4(3-carboxy-propionyl amino)-2-methyl-pentanoic acid or a pharmaceutically acceptable salt thereof; and

wherein said components (i) and (ii) are administered in one unit dose form or in two separate unit dose forms.

2. The method of claim **1** wherein components (i) and (ii) are administered in one unit dose form.

3. The method of claim **1** wherein components (i) and (ii) are administered separately in two separate unit dose forms.

4. The method of claim **1** wherein the condition or disease is hypertension.

5. The method of claim **1** wherein the condition or disease is heart failure.

6. The method of claim **2**, wherein the one unit dosage form is for oral administration.

7. The method of claim **2**, wherein the one unit dosage form is a tablet or capsule.

8. The method of claim **7**, wherein the tablet is a coated tablet.

9. The method of claim **3**, wherein the two separate unit dose forms are for oral administration.

10. The method of claim **3**, wherein the two separate unit dose forms are in a kit.

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